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INTERNATIONAL PRELIMINARY REPORT ON PATEN (Chapter II of the Patent Cooperation Treaty)

(PCT Article 36 and Rule 70)

	OR FURTHER ACTION See For	rm PCT/IPEA/416		
PU0407-PCT	ernational filing date (day/month/year)	Priority date (day/month/year)		
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International Patent Classification (IPC) or national classification and IPC				
See Supplemental Box				
Applicant				
GE HEALTHCARE BIO-SCIEN	CES AB et al			
		w this International Preliminary Examining		
This report is the international prelim: Authority under Article 35 and transi	mitted to the applicant according to Ar			
2. This REPORT consists of a total of 6 sheets, including this cover sheet.				
3. This report is also accompanied by ANNEXES, comprising:				
cont to the applicant and	a. (sent to the applicant and to the International Bureau) a total of sheets, as follows:			
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and/or sheets cor	ntaining rectifications authorized by the	is Authority (see Rule 70.16 and Section 607 of the		
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b. (sent to the International Bureau only) a total of (indicate type and number of electronic carrier(s))				
, containing a sequence listing and/or tables related thereto, in electronic form only, as indicated in the Supplemental Box Relating to Sequence Listing (see Section 802 of the				
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4. This report contains indications related Box No. I Basis of the				
Box No. II Priority Box No. III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability				
Box No. IV Lack of unity of invention				
Box No. V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability, citations and explanations supporting such statement				
Box No. VI Certain documents cited				
Box No. VII Certain defects in the international application				
Box No. VII Certain observations on the international application				
DONATO, VALL				
Date of submission of the demand	Date of comp	pletion of this report		
01-09-2005	29-05-2	2006		
Name and mailing address of the IPEA/SE	Authorized of	officer		
Patent- och registreringsverket				
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Form PCT/IPEA/409 (cover sheet) (April 2005)

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

International application No.

INTERNATIONAL PRELIMINARY REPORT ON PATENTABLETT	PCT/SE2005/000229
Supplemental Box	
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International patent classification (IPC)	
C12N15/10(2006.01) B01D 15/08 (2006.01)	

Form PCT/IPEA/409 (Supplemental Box) (April 2005)

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

International application No.

PCT/SE2005/000229

1. With regard to the language, this report is based on: the international application in the language in which it was filted a translation of the international application into which is the language of a translation furnished for the purposes of: international search (Rules 12.3(a) and 23.1(b)) publication of the international application (Rule 12.4(a)) international preliminary examination (Rules 53.2(a)) 2. With regard to the elements of the international application, this report is based on (replacement sheets which have been furnished to the receiving Office in responses to an invitation under Article 14 are referred to in this report as "originally filted" and are not annexed to this report): the international application as originally filed/furnished the description: pages pages* received by this Authority on pages* received by this Authority on the claims: pages pages* received by this Authority on pages* received by this Authority on the drawings: pages pages* received by this Authority on as originally filed/furnished received by this Authority on pages* received by this Authority on as originally filed/furnished pages* received by this Authority on pages* received by this Authority on as originally filed/furnished pages* received by this Authority on pages* received by this Authority on pages* received by this Authority on as originally filed/furnished pages* received by this Authority on as originally filed/furnished pages* received by this Authority on as originally filed/furnished pages* received by this Authority on as originally filed/furnished pages* received by this Authority on as originally filed/furnished pages* received by this Authority on pages* received by this Authority on as originally filed/furnished pages* received by this Authority on pages* received by this Authority on as originally filed/furnished pages* received by this Authority on pages* received by this Authority on as originally filed/furnished pages* received by this Authority on as originally filed/furnished pages* rec	Box	No. I	Basis of the report
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* If item 4 applies, some or all of those sheets may be marked "superseded."			

International application No.

PCT/SE2005/000229

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; Box No. V citations and explanations supporting such statement 1. Statement YES Novelty (N) Claims 7-8.16 NO Claims 1.2.11.12 YES Inventive step (IS) Claims Claims 1-6.9-15.17-22 YES Industrial applicability (IA) Claims 1-22 NO Claims

2. Citations and explanations (Rule 70.7)

The invention relates to methods for the isolation of plasmids using a separation matrix with anion exchange groups. chosen pore size distribution does not allow access of plasmids to the pore surfaces.

The most relevant documents cited in the International Search Report are:

D1: WO9963076A1 D2: WO0137987A1 D3: US6270970B1

Document D1 discloses a method of purifying plasmids using a TMAE anion exchange chromatographic column (see claims 1-3). The used matrix is a fractogel TMAE anion exchange resin. These resins are known to have particle sizes between 20-40 μm for TMAE S and 40-90 µm for TMAE M. The pore size is about 800 Å (see Merck website).

Thus, D1 is considered to disclose a method of isolating plasmids with the steps of providing a separation matrix comprised of porous carriers, which carrier present anion exchange groups on external surfaces as well as pore surfaces and a pore size distribution that does not allow access of plasmids to pore surfaces; contacting said matrix with a liquid to absorb plasmids to ligands present on the external surfaces of the separation matrix

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Supplemental Box

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Document D2 discloses separation methods for plasmids. In example 3 a separation of plasmids is performed with anion exchange chromatography. The plasmids are bound to the separation medium B and its charged outer surfaces of the anion-exchanger. It is not stated that the plasmids have access to the pores.

Document D3 relates to mixed-bed solid phases for isolation of nucleic acids such as plasmids. The solid phase of the different beds comprise magnetic silica particles (particle size below 15 μ m), see column 12. The solid phase can be with or without pores with size sufficiently large to admit the target nucleic acid in to the interior of the particles. The anion exchanger phase can be Sepharose but is not limited thereto.

With background of D1-D3, and as a consequence of unclear claims (see box VIII), the method according to claim 1 and the use according to claim 11 lacks novelty. Further, the DNA exclusion limits covered by D1-D3 are assumed to be at least about 270 base pairs. Therefore, also claims 2 and 12 lacks novelty.

The claims 3-6, 9-10, 13-15 and 17 are considered to involve particular detail executions obvious to a person skilled in the art. Therefore, the invention according to these claims is not considered to involve an inventive step.

It is also considered to be obvious to a person skilled in the art to develop a kit for the method described in D1 or D2. Therefore the invention according to claims 18-22 lacks an inventive step.

Claims 7-8 and 16 are novel and considered to involve an inventive step.

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Box No. VIII Certain observations on the international application

The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made:

Claims 1, 11 and 18 do not meet the requirements of Article 6 PCT in that the matter for which protection is sought is not clearly defined. The claims attempts to define the subject-matter in terms of the result to be achieved (...pore size distribution that does not allow access of plasmids...) which merely amounts to a statement of the underlying problem. The technical features necessary for achieving this result should be added.

In claims 2, 12 and 19 the matrix is characterised by a DNA exclusion limit of at least about 270 base pairs. This way of characterising a matrix is known in the field but is not a common way of defining and comparing gels. Further, the limit of "about" 270 base pairs is unclear (see PCT GL 5.38).

Claims 1-2, 11-12 and 18-19 have been drafted as separate independent claims of the same category. They appear to relate effectively to the same subject-matter and to differ from each other only with regard to the choice of specific words. The aforementioned claims therefore lack conciseness. See PCT Article 6 and 5.42 Guidelines.